



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
2098 Gaither Road  
Rockville MD 20850

NOV 23 2004

Dr. Pauline Armstrong  
Regulatory Affairs  
Randox Laboratories, Ltd.  
55 Diamond Road  
Crumlin, Co. Antrim  
United Kingdom BT29 4QY

Re: k041142  
Trade/Device Name: evidence® Cannabinoids Assay and  
evidence® Drugs of Abuse Calibrators  
Regulation Number: 21 CFR 862.3870  
Regulation Name: Cannabinoid test system  
Regulatory Class: Class II  
Product Code: LDJ, DKB  
Dated: October 26, 2004  
Received: October 28, 2004

Dear Dr. Armstrong:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. In addition, FDA may publish further announcements concerning your device in the Federal Register.

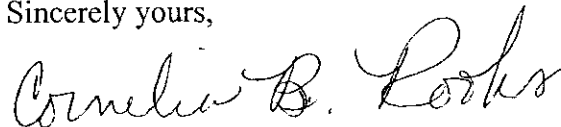
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Parts 801 and 809); and good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820).

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This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific information about the application of labeling requirements to your device, or questions on the promotion and advertising of your device, please contact the Office of In Vitro Diagnostic Device Evaluation and Safety at (301) 594-3084. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>.

Sincerely yours,

A handwritten signature in black ink, reading "Cornelia B. Rooks". The signature is fluid and cursive, with the first name "Cornelia" and middle initial "B." written in a smaller, more compact script, and the last name "Rooks" written in a larger, more prominent cursive style.

Cornelia B. Rooks, MA  
Acting Director  
Division of Chemistry and Toxicology  
Office of In Vitro Diagnostic Device  
Evaluation and Safety  
Center for Devices and  
Radiological Health

Enclosure

## Indications for Use

510(k) Number (if known): \_\_\_\_\_

Device Name: evidence® CANNABINOIDS ASSAY AND  
evidence® DRUGS OF ABUSE CALIBRATORS

### Indications For Use:

The **evidence®** Cannabinoids test has been designed for use only on the **evidence®** analyser to detect cannabinoids in urine, using a cut-off of 50ng/ml. Qualitative results obtained can be utilised in the diagnosis of cannabinoid use or abuse.

**This assay provides only a preliminary analytical test result which should be confirmed by a more specific method, such as GC/MS.**

The Cannabinoids Assay must only be used by suitably qualified laboratory personnel under appropriate laboratory conditions.

### The **evidence®** Drugs of Abuse Calibrators.

The **evidence®** Drugs of Abuse Calibrators are liquid Calibrators containing benzoylecgonine, amphetamine, methamphetamine, methadone and 11-nor-D9-THC-9 carboxylic acid.

There are 9 levels of calibrator. They have been developed for use in calibration of the **evidence®** system.

The **evidence®** Drugs of Abuse Calibrators must only be used by suitably qualified laboratory personnel under appropriate laboratory conditions.

Prescription Use ☒   
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use \_\_\_\_\_   
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRI, Office of In Vitro Diagnostic Devices (OIVD)

  
Division Sign-Off

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Office of In Vitro Diagnostic  
Device Evaluation and Safety

510(k)

K041142